

In the Claims

1-30 (canceled).

31 (currently amended). A vaccine composition that induces a protective an immune response against two or more subtypes of FIV in an animal susceptible to infection by FIV, comprising an effective amount of an FIV immunogen to induce said immune response, wherein said FIV immunogen comprises an immunogen or immunogens from or comprising at least two different FIV subtypes.

32 (previously presented). The vaccine composition according to claim 31, wherein said immunogen is or immunogens are, independently, selected from the group consisting of synthetic FIV peptide, natural or recombinant FIV protein or an immunogenic fragment of said FIV protein, cell-free whole or partial FIV virus, and a cell infected with FIV virus.

33 (previously presented). The vaccine composition according to claim 32, wherein said FIV virus or FIV-infected cell is treated in a manner to inactivate said FIV virus or the FIV virus infecting said cell prior to administration of said vaccine to said animal.

34 (previously presented). The vaccine composition according to claim 32, wherein said FIV virus or FIV-infected cell is treated in a manner to attenuate said FIV virus or the FIV virus infecting said cell prior to administration of said vaccine to said animal.

35 (previously presented). The vaccine composition according to claim 31, wherein said at least two different FIV subtypes are selected from the group consisting of subtypes A, B, C, and D.

36 (previously presented). The vaccine composition according to claim 31, wherein said at least two different FIV subtypes are subtypes A and D.

37 (previously presented). The vaccine composition according to claim 31, wherein said FIV immunogen is or immunogens are from an FIV virus strain selected from the group consisting of FIV<sub>Dix</sub>, FIV<sub>UK8</sub>, FIV<sub>Bang</sub>, FIV<sub>Aom1</sub>, FIV<sub>Aom2</sub>, FIV<sub>Pet</sub>, and FIV<sub>Shi</sub>.

38 (previously presented). The vaccine composition according to claim 32, wherein said cell is from the cell line designated FeT-1C having ATCC accession number CRL 11968 infected with FIV.

39 (previously presented). The vaccine composition according to claim 32, wherein said cell is from the cell line designated FeT-J having ATCC accession number CRL 11967 infected with FIV.

40 (previously presented). The vaccine composition according to claim 32, wherein said cell is from the cell line designated FL-4 having ATCC accession number CRL 10772 infected with FIV.

41 (previously presented). The vaccine composition according to claim 32, wherein said cell is from the cell line designated FeT-1M having ATCC accession number CRL 10775 infected with FIV.

42 (previously presented). The vaccine composition according to claim 32, wherein said cell is infected with FIV<sub>Shi</sub> and said cell is from a cell line designated Shi/FeT-1C having ATCC accession number CRL 11976.

43 (previously presented). The vaccine composition according to claim 32, wherein said cell is infected with FIV<sub>Bang</sub> and said cell is from a cell line designated Bang/FeT-J having ATCC accession number CRL 11975.

44-49 (canceled).

50 (previously presented). The vaccine composition according to claim 31, wherein said animal is a cat.

51 (previously presented). The vaccine composition according to claim 32, wherein said FIV protein comprises FIV envelope glycoprotein, or an immunogenic fragment thereof.

52 (previously presented). The vaccine composition according to claim 51, wherein said FIV envelope glycoprotein comprises the amino acid sequence shown in SEQ ID NO. 1.

53 (previously presented). The vaccine composition according to claim 32, wherein said FIV protein is a chimeric protein comprising amino acid sequences of a protein from at least two different FIV subtypes.

54 (previously presented). The vaccine composition according to claim 53, wherein said chimeric protein comprises FIV envelope glycoprotein.

55 (previously presented). The vaccine composition according to claim 31, wherein said vaccine composition further comprises an adjuvant.

56 (previously presented). The vaccine composition according to claim 55, wherein said adjuvant is selected from the group consisting of threonyl muramyl dipeptide, alum, complete Freund's, and incomplete Freund's.

57 (previously presented). The vaccine composition according to claim 31, wherein said vaccine composition is administered parenterally, orally, or nasally.

58 (previously presented). The vaccine composition according to claim 57, wherein said parenteral administration is by subcutaneous, intraperitoneal, or intramuscular injection.

59 (previously presented). The vaccine composition according to claim 32, wherein said FIV-infected cell is present in a dose of from about  $10^6$  cells to about  $10^8$  cells.

60 (previously presented). The vaccine composition according to claim 32, wherein said FIV-infected cell is present in a dose of from about  $5 \times 10^6$  cells to about  $7.5 \times 10^7$  cells.

61 (previously presented). The vaccine composition according to claim 32, wherein said cell-free whole or partial FIV virus is present in a dose from about 0.1 mg to about 5 mg.

62 (previously presented). The vaccine composition according to claim 32, wherein said cell-free whole or partial FIV virus is present in a dose from about 0.2 mg to about 2 mg.

63 (previously presented). The vaccine composition according to claim 31, wherein said FIV immunogen comprises (a) cells infected with FIV of a first subtype and (b) cell-free whole or partial FIV of a second subtype, wherein said first and second subtype of said FIV are selected from the group consisting of A, B, C, and D, and wherein said first and second subtype of FIV are not the same.

64-107 (canceled).